FDA 510(k) Premarket Notification

K123805 E-Z CHEK BLOOD LEAK TEST STRIPS

CLEARED

Mar 20, 2013
99 days to decision

K123805 · Product code: **FJD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k123805/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector, Leak, Blood (FJD)
Date received	Dec 11, 2012
Decision date	Mar 20, 2013
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Reprocessing Products Corp**
Location	Tucson, AZ, US
Contact	TED WILLIAMS
510(k) history	6 submissions · 6 cleared · 1995-2013